AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Previously Presented) A compound of the general formula (I)

wherein

A represents an aryl or heteroaryl ring,

 R^1 , R^2 and R^3 independently from each other represent hydrogen, halogen, nitro, cyano, C_1 - C_6 -alkyl, hydroxy or C_1 - C_6 -alkoxy, wherein C_1 - C_6 -alkyl and C_1 - C_6 -alkoxy can be further substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy and C_1 - C_4 -alkoxy,

 $R^4 \qquad \text{represents trifluoromethylcarbonyl, C_1-C_6-alkylcarbonyl, C_1-C_6-alkoxycarbonyl, hydroxycarbonyl, aminocarbonyl, mono- or di-C_1-C_4-alkylaminocarbonyl, C_6-C_{10}-arylaminocarbonyl, arylcarbonyl, heteroarylcarbonyl, heterocyclylcarbonyl, heterocyclyl or cyano, wherein C_1-C_6-alkylcarbonyl, C_1-C_6-alkoxycarbonyl, mono- and di-C_1-C_4-alkylaminocarbonyl can be further substituted with one to three identical or different radicals selected$

from the group consisting of C₃-C₈-cycloalkyl, hydroxy, C₁-C₄-alkoxy, C₁-C₄alkoxycarbonyl, hydroxycarbonyl, aminocarbonyl, mono- and di-C₁-C₄alkylaminocarbonyl, C₁-C₄-alkylcarbonylamino, (C₁-C₄-alkylcarbonyl)-C₁-C₄alkylamino, cyano, amino, mono- and di-C₁-C₄-alkylamino, heteroaryl, heterocyclyl and tri-(C₁-C₆-alkyl)-silyl, and wherein heteroarylcarbonyl, heterocyclylcarbonyl, heteroaryl and heterocyclyl can be further substituted with C₁-C₄alkyl,

 R^5 represents C₁-C₄-alkyl, which can be substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy, C₁-C₆alkoxy, C₁-C₆-alkenoxy, C₁-C₆-alkylthio, amino, mono- and di-C₁-C₆-alkylamino, arylamino, hydroxycarbonyl, C₁-C₆-alkoxycarbonyl and the radical -O-C₁-C₄alkyl-O- C_1 - C_4 -alkyl,

or

- R^5 represents amino,
- R^6 represents hydrogen, C₁-C₆-alkyl, formyl, aminocarbonyl, mono- or di-C₁-C₄alkylaminocarbonyl, C₃-C₈-cycloalkylcarbonyl, C₁-C₆-alkylcarbonyl, C₁-C₆alkoxycarbonyl, N-(C₁-C₄-alkylsulfonyl)-aminocarbonyl, N-(C₁-C₄-alkylsulfonyl)-N-(C₁-C₄-alkyl)-aminocarbonyl, heteroaryl, heteroaryl, carbonyl or heterocyclylcarbonyl, wherein C₁-C₆-alkyl, mono- and di-C₁-C₄alkylaminocarbonyl, C₁-C₆-alkylcarbonyl, C₁-C₆-alkoxycarbonyl, heteroaryl and heterocyclyl can be substituted with one to three identical or different radicals selected from the group consisting of aryl, heteroaryl, hydroxy, C_1 - C_4 -alkoxy, hydroxycarbonyl, C₁-C₆-alkoxycarbonyl, aminocarbonyl, mono- and di-C₁-C₄alkylaminocarbonyl, amino, mono- and di-C₁-C₄-alkylamino, C₁-C₄alkylcarbonylamino, tri-(C₁-C₆-alkyl)-silyl, cyano, mono- and di-C₁-C₄-

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alkylamino- C_1 - C_4 -alkylaminocarbonyl, C_1 - C_4 -alkoxy- C_1 - C_4 -alkylaminocarbonyl and halogen,

or

R⁶ represents a moiety of the formula

wherein

 R^{6A} is selected from the group consisting of hydrogen and C_1 - C_6 -alkyl, and

n represents an integer of 1 or 2,

 R^7 represents halogen, nitro, cyano, C_1 - C_6 -alkyl, hydroxy or C_1 - C_6 -alkoxy, wherein C_1 - C_6 -alkyl is further substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy and C_1 - C_4 -alkoxy, and C_1 - C_6 -alkoxy can be further substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy and C_1 - C_4 -alkoxy,

and

Y¹, Y², Y³, Y⁴ and Y⁵ independently from each other represent CH or N, wherein the ring contains either 0, 1 or 2 nitrogen atoms,

or a pharmaceutically acceptable salt thereof.

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2. (Previously Presented) The compound of general formula (I) according to Claim 1, wherein

- A represents an aryl or heteroaryl ring,
- R^1 , R^2 and R^3 independently from each other represent hydrogen, halogen, nitro, cyano, C_1 - C_6 -alkyl, hydroxy or C_1 - C_6 -alkoxy, wherein C_1 - C_6 -alkyl and C_1 - C_6 -alkoxy can be further substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy and C_1 - C_4 -alkoxy,
- represents C_1 - C_6 -alkylcarbonyl, C_1 - C_6 -alkoxycarbonyl, C_1 - C_6 -alkenoxycarbonyl, hydroxycarbonyl, aminocarbonyl, mono- or di- C_1 - C_4 -alkylaminocarbonyl, C_6 - C_{10} -arylaminocarbonyl, heteroarylcarbonyl, heterocyclylcarbonyl, heteroaryl, heterocyclyl or cyano, wherein C_1 - C_6 -alkylcarbonyl, C_1 - C_6 -alkoxycarbonyl, mono- and di- C_1 - C_4 -alkylaminocarbonyl can be further substituted with one to three identical or different radicals selected from the group consisting of C_3 - C_8 -cycloalkyl, hydroxy, C_1 - C_4 -alkoxy, C_1 - C_4 -alkoxycarbonyl, hydroxycarbonyl, aminocarbonyl, mono- and di- C_1 - C_4 -alkylaminocarbonyl, C_1 - C_4 -alkylcarbonyl-amino, amino, mono- and di- C_1 - C_4 -alkylamino, heteroaryl, heterocyclyl and tri- $(C_1$ - C_6 -alkyl)-silyl,
- R^5 represents C_1 - C_4 -alkyl, which can be substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy, C_1 - C_6 -alkoxy, C_1 - C_6 -alkenoxy, C_1 - C_6 -alkylthio, amino, mono- and di- C_1 - C_6 -alkylamino, arylamino, hydroxycarbonyl, C_1 - C_6 -alkoxycarbonyl and the radical -O- C_1 - C_4 -alkyl,

or

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R⁵ represents amino,

or

R⁶ represents a moiety of the formula

$*$
 $^{\circ}$ $^{\circ}$

wherein

 R^{6A} is selected from the group consisting of hydrogen and $C_1\text{-}C_6\text{-alkyl}$, and

n represents an integer of 1 or 2,

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 R^7 represents halogen, nitro, cyano, C_1 - C_6 -alkyl, hydroxy or C_1 - C_6 -alkoxy, wherein C_1 - C_6 -alkyl is further substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy and C_1 - C_4 -alkoxy, and C_1 - C_6 -alkoxy can be further substituted with one to three identical or different radicals selected from the group consisting of halogen, hydroxy and C_1 - C_4 -alkoxy,

and

 Y^1 , Y^2 , Y^3 , Y^4 and Y^5 independently from each other represent CH or N, wherein the ring contains either 0, 1 or 2 nitrogen atoms.

- 3. (Previously Presented) The compound of general formula (I) according to Claim 1, wherein
 - A represents a phenyl, naphthyl or pyridyl ring,
 - R¹, R² and R³ independently from each other represent hydrogen, fluoro, chloro, bromo, nitro, cyano, methyl, ethyl, trifluoromethyl or trifluoromethoxy,
 - R^4 represents C_1 - C_6 -alkylcarbonyl, C_1 - C_6 -alkoxycarbonyl, hydroxycarbonyl, aminocarbonyl, mono- C_1 - C_4 -alkylaminocarbonyl or cyano, wherein C_1 - C_6 -alkylcarbonyl, C_1 - C_6 -alkoxycarbonyl and mono- C_1 - C_4 -alkylaminocarbonyl can be substituted with one to three identical or different radicals selected from the group consisting of C_3 - C_8 -cycloalkyl, hydroxy, C_1 - C_4 -alkoxy, C_1 - C_4 -alkoxycarbonyl, amino, mono- or di- C_1 - C_4 -alkylamino, heteroaryl and heterocyclyl,
 - R⁵ represents methyl or ethyl,

 R^6 represents hydrogen, C_1 - C_6 -alkyl, mono- or di- C_1 - C_4 -alkylaminocarbonyl, C_1 - C_6 -alkylcarbonyl, C_1 - C_6 -alkoxycarbonyl or heterocyclylcarbonyl, wherein C_1 - C_6 -alkyl and C_1 - C_6 -alkoxycarbonyl can be substituted with one to three identical or different radicals selected from the group consisting of heteroaryl, hydroxy, C_1 - C_4 -alkoxy, hydroxycarbonyl, C_1 - C_6 -alkoxycarbonyl, aminocarbonyl, mono- and di- C_1 - C_4 -alkylaminocarbonyl, cyano, amino, mono- and di- C_1 - C_4 -alkylamino,

or

R⁶ represents a moiety of the formula

wherein

 R^{6A} is selected from the group consisting of hydrogen and $C_1\text{-}C_4\text{-alkyl}$, and

- n represents an integer of 1 or 2,
- R⁷ represents halogen, nitro, cyano, trifluoromethyl, or trifluoromethoxy,

and

 Y^1 , Y^2 , Y^3 , Y^4 and Y^5 each represent CH.

4. (Previously Presented) The compound of general formula (I) according to Claim 1, wherein

A represents a phenyl or a pyridyl ring,

R¹ and R³ each represent hydrogen,

- R² represents fluoro, chloro, bromo, nitro or cyano,
- R^4 represents cyano, C_1 - C_4 -alkylcarbonyl or C_1 - C_4 -alkoxycarbonyl, wherein C_1 - C_4 -alkoxycarbonyl can be substituted with a radical selected from the group consisting of hydroxy, C_1 - C_4 -alkoxy, C_1 - C_4 -alkoxycarbonyl, mono- and di- C_1 - C_4 -alkylamino, heteroaryl and heterocyclyl,
- R⁵ represents methyl,
- R^6 represents hydrogen, C_1 - C_4 -alkyl, mono- or di- C_1 - C_4 -alkylaminocarbonyl, C_1 - C_4 -alkylcarbonyl or C_1 - C_4 -alkoxycarbonyl, wherein C_1 - C_4 -alkyl and C_1 - C_4 -alkoxycarbonyl can be substituted with a radical selected from the group consisting of heteroaryl, hydroxy, C_1 - C_4 -alkoxy, hydroxycarbonyl, aminocarbonyl, mono- and di- C_1 - C_4 -alkylaminocarbonyl, amino, mono- and di- C_1 - C_4 -alkylamino,

or

R⁶ represents a moiety of the formula

wherein

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R^{6A} is selected from the group consisting of hydrogen and methyl,

R⁷ represents trifluoromethyl or nitro,

and

 Y^1 , Y^2 , Y^3 , Y^4 and Y^5 each represent CH.

- 5. (Previously presented) The compound of general formula (I) according to claim 1, wherein A is phenyl or pyridyl.
- 6. (Previously Presented) The compound of general formula (I) according to claim 1, wherein R¹ is hydrogen.
- 7. (Previously Presented) The compound of general formula (I) according to claim 1, wherein \mathbb{R}^2 is cyano.
- 8. (Previously Presented) The compound of general formula (I) according to claim 1, wherein R³ is hydrogen.
- 9. (Previously Presented) The compound of general formula (I) according to claim 1, wherein R^4 is C_1 - C_4 -alkoxycarbonyl optionally substituted by hydroxy or wherein R^4 is C_1 - C_4 -alkylcarbonyl.
- 10. (Previously Presented) The compound of general formula (I) according to claim 1, wherein R⁵ is methyl.
- 11. (Previously Presented) The compound of general formula (I) according to claim 1, wherein \mathbb{R}^6 is hydrogen.

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12. (Previously Presented) The compound of general formula (I) according to claim 1, wherein \mathbb{R}^7 is trifluoromethyl or nitro.

13. (Previously Presented) A compound of general formula (IA)

$$R^{1}$$
 R^{4}
 R^{4}
 R^{6}
 R^{3}
 CF_{3}

wherein

Z represents CH or N, and

 R^1 , R^3 , R^4 and R^6 have the meaning indicated in claim 1.

14. (Previously Presented) A process for synthesizing the compounds of general formula (I), as defined in claim 1 by condensing compounds of general formula (II)

$$R^{1}$$
 A
 CHO
(II),

wherein

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A, R¹ and R² have the meaning indicated in claim 1,

with compounds of general formula (III)

wherein

R⁴ and R⁵ have the meaning indicated in claim 1,

and compounds of general formula (IV)

$$\begin{array}{c}
NH_2 \\
NH_2 \\
O \\
Y_1^1 \longrightarrow Y^5 \\
Y_2^2 \longrightarrow Y^3 \longrightarrow Y^4
\end{array}$$
(IV),

wherein

 R^3 , R^7 , and Y^1 to Y^5 have the meaning indicated in claim 1,

in the presence of an acid either in a three-component / one-step reaction or sequentially to give compounds of the general formula (IB)

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$$R^{1}$$
 A
 R^{4}
 NH
 R^{5}
 N
 O
 Y_{1}^{1}
 Y_{2}^{5}
 Y_{3}^{7}
 Y_{4}^{7}
 X_{1}^{7}
 X_{2}^{7}
 X_{3}^{7}
 X_{4}^{7}

wherein

A, R^1 to R^5 , R^7 , and Y^1 to Y^5 have the meaning indicated in claim 1,

optionally followed by reaction of the compounds of general formula (IB) with compounds of the general formula (V)

$$R^{6*}-X$$
 (V),

wherein

 R^{6*} has the meaning of R^6 as indicated in claim 1, but does not represent hydrogen, and X represents a leaving group,

in the presence of a base.

- 15. (Previously Presented) A composition containing at least one compound of general formula (I) as defined in claim 1 and a pharmacologically acceptable diluent.
- 16. (Canceled)
- 17. (Previously Presented) A process for preparation of a composition, said process comprising a step of bringing the compounds of general formula (I) as defined in claim 1

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together with customary auxiliaries into a suitable application form; wherein said composition contains at least one compound of general formula (I) and a pharmacologically acceptable diluent.

- 18. (Canceled)
- 19. (Currently Amended) A method of treating acute and chronic inflammatory, ischaemic or remodelling processes chronic obstructive pulmonary disease or acute myocardial infarction, said method comprising administering a therapeutically effective amount of a compound of claim 1.
- 20. (Canceled)
- 21. (Previously Presented) The method of claim 19, wherein a neutrophil elastase inhibitory amount is administered.
- 22. (Previously Presented) A composition containing at least one compound of general formula (IA) as defined in claim 13 and a pharmacologically acceptable diluent.
- 23. (Previously Presented) A process for preparation of a composition, said process comprising a step of bringing the compounds of general formula (IA) as defined in claim 13 together with customary auxiliaries into a suitable application form; wherein said composition contains at least one compound of general formula (IA) and a pharmacologically acceptable diluent.
- 24. (Previously Presented) Ethyl 4-(4-cyanophenyl)-6-methyl-1-(3-methylphenyl)-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxylate, or a pharmaceutically acceptable salt thereof.